

## HYDROGENOLYSIS OF SUGAR FEEDSTOCK

The present invention relates to a homogeneous process for the production of glycols from sugar derived feedstocks. More particularly, it relates to a homogeneous hydrogenolysis process which can be carried out in the presence of water. Most particularly it relates to a homogeneous hydrogenolysis process for a feedstock comprising one or more of polyols, alditols, aldoses, polymers of aldoses and starch.

For ease of reference the feedstock comprising one or more of polyols, alditols, aldoses, polymers of aldoses such as starch and cellulose will be described generally as a "sugar feedstock". The polymers of aldoses include homopolymers and copolymers.

Many catalyst systems are known which are suitable for use in the hydrogenolysis of sugars. Traditionally such reactions are carried out using heterogenous catalysts and often high temperature and pressures. Typically temperatures in the range of about 200°C to about 275°C are required with pressures in the region of from about 1000 psig to about 4000 psig. Many of these require the use of basic promoters to prevent catalyst degradation and/or to promote catalyst activity. However, the use of these promoters adds significantly to the cost of the reaction. The use of sulphur containing additives have been suggested to increase the selectivity of the catalyst. However, this increase in selectivity is often at the expense of a loss of activity. Examples of heterogeneous processes can be found in US6479713, US6291725, US5326912, US5354914, US5600028, US5403805, US5210335, US5107018, US5107018, FR2603276, US4496780, US4476331, US443184, US4401823, US4380678, US4404411, US4366332, GB988040, US3011002, US282603, GB490211, GB430576, Abreau et al, Biomass and Bioenergy 9, 587 (1995) and J.Catalysis 208 248 (2002) Fabre et al.

Homogeneous process have also been suggested and examples of these can be found in US5118883, US5026927, US3935284, US6080898, US4642394, US5097089, US3454644, J.Organomet. Chem. 417 41 (1991) G Braca et al, J. Molecular Catal. 22 138 (1983) and J. Molecular Catal. 16 349 (1982).

Whilst some of these processes go some way to providing a commercial process, they suffer from certain disadvantages and drawbacks. In particular, they are costly to operate, many require the presence of a strong basic promoter and are temperature sensitive. For example, the process of US 5026927 operates at a temperature of from 75°C to about 150°C and that of US 3935284 requires a temperature of 150°C or less. It is stated in US 3935284 that at temperatures in excess of 150°C, decarbonylation occurs to produce a carbonyl-ruthenium species which is a less active catalyst.

It is therefore desirable to provide a process which provides a cost-effective process for sugar hydrogenolysis and which utilises a catalyst regime that has the required levels of selectivity and activity.

Thus according to the present invention there is provided a process for the hydrogenolysis of a sugar feedstock in the presence of a catalyst comprising:

(a) ruthenium or osmium; and

(b) an organic phosphine;

and wherein the hydrogenolysis is carried out in the presence of water and at a temperature of greater than 150°C.

By "homogeneous process" we mean that the catalyst is dissolved in the solvent for the reaction and that at least some of the water present and at least some of the sugar feedstock must be in phase with the catalyst. Where excess water and/or excess feedstock is present, the excess may form a separate phase to that comprising the catalyst. Additionally, or alternatively, the product may form a separate phase.

As detailed above, the sugar feedstock may be a feedstock comprising one or more of polyols, alditols, aldoses and polymers of aldoses such as cellulose and starch. Examples of alditols and aldoses suitable for use in the process of the present invention include those having from C<sub>3</sub> to C<sub>12</sub>, more particularly C<sub>3</sub> to C<sub>6</sub>. Examples of suitable feedstocks include glucose, sucrose, xylose, arabinose, mannose, mannitol, sorbitol, xylitol, arabinol, glycerol and mixtures thereof. The sugar feedstock may be provided from natural or synthetic sources or mixtures thereof.

Where the sugar feedstock is water soluble, the water may be present as the solvent for the reaction. Alternatively, a solvent may be used. Where a solvent is used, the water will be present as an additive in the solvent. In another alternative arrangement, the sugar feedstock or the product of the reaction may be the solvent. In one arrangement at least 1% by weight of water is present.

Where the sugar feedstock is non-water soluble or has low water-solubility, such as for example a sugar having a higher carbon content such as high molecular weight polymeric alditols, the feedstock or product may be the solvent for the reaction or an organic solvent may be used and the water may be present as an additive. In this case, it may be present in the solvent in any suitable amount and preferably in an amount of from about 1% up to the solubility limit of the water in the solvent. Additional water may be present in a separate aqueous phase.

The process of the present invention provides a method for the hydrogenolysis of sugars which can be carried out at higher temperatures than has been achievable heretofore to increase activity while maintaining the desired level of selectivity.

Further, it has been found that the presence of water is beneficial in terms of catalyst stability. It is noted that in prior art systems, decarbonylation is noted and the carbon monoxide formed is said to strongly inhibit the catalyst. Without wishing to be bound by any theory, it is believed that the presence of water allows a side reaction to occur in the hydrogenation reactor in which any carbon monoxide produced reacts with the water to form carbon dioxide and hydrogen via the water gas shift reaction. This carbon dioxide and hydrogen may be further reacted to form methane. These gases can be readily removed from the reaction system. It will therefore be appreciated that the need to provide a separate methanation unit in the recycling system for vent gases is obviated.

A further advantage of the present invention is that the removal of the carbon monoxide as detailed above allows for effective regeneration of the catalyst. Thus the process offers extended catalyst life which in turn improves the economics of the reaction.

As detailed above, where the sugar feedstock is soluble in water, the water may act as the solvent. However, the method of the present invention may be conducted in the absence of a solvent, i.e. the starting material or reaction product may be a solvent for the reaction. However, if a solvent is used, any suitable solvent may be selected and examples of suitable solvents include, but are not limited to tetrahydrofuran, tetraethyleneglycol dimethyl ether, N-methyl pyrrolidone, diethyl ether, ethyleneglycol dimethylether, dioxane, 2-propanol, 2-butanol, secondary alcohols, tertiary alcohols, lactams and N-methyl caprolactam.

The catalyst of the present invention is a ruthenium/phosphine or osmium/phosphine catalyst with a ruthenium/phosphine catalyst being particularly preferred. The ruthenium is generally provided as a ruthenium compound although halides are not preferred. Suitable compounds are those which can be converted to active species under the reaction conditions and include nitrates, sulphates, carboxylates, beta diketones, and carbonyls. Ruthenium oxide, carbonyl ruthenates and complex compounds of ruthenium, including hydridophosphineruthenium complexes, may also be used. Specific examples include, but are not limited to, ruthenium nitrate, ruthenium dioxide, ruthenium tetraoxide, ruthenium dihydroxide, ruthenium acetylacetonate, ruthenium acetate, ruthenium maleate, ruthenium succinate, tris-(acetylacetonate)ruthenium, pentacarbonylruthenium, dipotassium tetracarbonyl-ruthenium, cyclo-pentadienyldicarbonyltriruthenium, ruthenium dihydroxide, bis(tri-n-butylphosphine)tricarbonylruthenium, dodecacarbonyltriruthenium, tetrahydride-decacarbonyltetraruthenium, and undecacarbonylhydridetetriruthenate. Corresponding compounds may be used where the catalyst is formed from osmium.

The catalyst may be preformed or generated in situ. Where an electron rich phosphine such as tris-1, 1, 1 - (diethylphosphinomethyl)ethane, is to be used it may be preferable to preform the catalyst in the absence of water prior to commencing the process of the present invention.

The ruthenium/osmium compound may be present in any suitable amount. However, it is preferably present in an amount of from 0.0001 to 5 mol, preferably 0.005 to 1 mol, as ruthenium/osmium per liter of reaction solution.

Any suitable phosphine may be used. Compounds which provide tridentate, bidentate and monodentate ligands may be used. Where the metal is ruthenium, tridentate phosphines are particularly preferred. Examples of suitable phosphine compounds include trialkylphosphines, dialkylphosphines, monoalkylphosphines, triarylphosphines, diarylphosphines, monoarylphosphines, diarylmonoalkyl phosphines and dialkylmonoaryl phosphines. Specific examples include but are not limited to tris-1,1,1-(diphenylphosphinomethyl)methane, tris-1,1,1-(diphenylphosphinomethyl)ethane, tris-1,1,1-(diphenylphosphinomethyl)propane, tris-1,1,1-(diphenylphosphino-methyl)butane, tris-1,1,1-(diphenylphosphinomethyl)-2,2dimethylpropane, tris-1,3,5-(diphenylphosphino-methyl)cyclohexane, tris-1,1,1-(dicyclohexylphosphinomethyl)ethane, tris-1,1,1-(dimethylphosphinomethyl)ethane, tris-1,1,1-(diethylphosphinomethyl)ethane, 1,5,9-triethyl-1,5,9-triphosphacyclododecane, 1,5,9-triphenyl-1,5,9-triphosphacyclododecane, bis(2-diphenylphosphinoethyl)phenylphosphine, bis-1,2-(diphenyl phosphino)ethane, bis-1,3-(diphenyl phosphino)propane, bis-1,4-(diphenyl phosphino)butane, bis-1,2-(dimethyl phosphino)ethane, bis-1,3-(diethyl phosphino)propane, bis-1,4-(dicyclohexyl phosphino)butane, tricyclohexylphosphine, trioctyl phosphine, trimethyl phosphine, tripyridyl phosphine, triphenylphosphine with tris-1,1,1-(diphenylphosphinomethyl)-ethane being particularly preferred. Particularly advantageous results are achieved with tridentate facially capped phosphines with tris-1,1,1- (diarylphosphinomethyl)alkane and tris-1,1,1-(dialkylphosphinomethyl)alkane being particularly preferred.

The phosphine compound may be present in any suitable amount. However, it is preferably present in an amount of from 0.0001 to 5 mol, preferably 0.005 to 1 mol, as phosphine per liter of reaction solution.

Whilst a strong base, such as potassium hydroxide, may be added they are not believed to have any significant benefit to the selectivity of the process. Examples of base additives include any of those identified in the prior art.

However, in one arrangement of the present invention an increase in selectivity may be noted where a second phosphine is present. The second phosphine will generally be a phosphine which is a more weakly coordinating ligand to the ruthenium or osmium than the first

phosphine compound. Examples of suitable second phosphines include triphenylphosphine and phosphine oxides such as triphenylphosphine oxide. Without wishing to be bound by any theory, these weakly co-ordinating ligands may compete with the active site at the metal thus preventing coordination of the product and thereby any undesirable side reactions from occurring. Alternatively, other weakly coordinating ligands such as amines may be used.

Any suitable reaction temperature in excess of 150°C may be used. However, in the process of the present invention, particular advantages may be noted if the hydrogenolysis is carried out at temperatures in the region of from about 190°C to about 260°C, more preferably 200°C to about 250°C.

Any suitable pressure may be used with a reaction pressure of from about 250 psig to about 2000 psig, being preferred. More preferably a pressure of from 800 psig to 1200 psig may be used and most preferably a pressure of about 1000 psig may be used. However, it will be understood that if a volatile solvent is used a higher reactor pressure may be desirable due to the high partial pressure of the solvent in the reactor.

The process may be carried out either in a batch system or in a continuous system. High intensity reactors such as intensive gas/liquid mixing reactors may be used. However, it will be understood that the process of the present invention is particularly suitable for use in a continuous system since the catalyst is not poisoned by carbon monoxide or if poisoning in this way occurs, the catalyst can be regenerated by reaction with the water.

Where the catalyst is removed from the reactor, for example, with a product removal stream, it may be recycled by any suitable means to the reactor. The catalyst may be separated from the product stream by any suitable means. Suitable means include extraction, distillation, gas stripping and membrane separation. In some circumstances, the catalyst may be immobilised on a support to assist the recovery. In this arrangement, the immobilised catalyst may be recovered by filtration.

A pre-reduction step may be included to improve the selectivity to the desired product. In one arrangement, the pre-reduction step may be carried out in the same reactor to the main reaction. In one alternative arrangement the pre-reduction may be carried out in a different reactor. Where the same reactor is used, the pre-reduction step may be carried out within different zones within the reactor or the same zone. Where the same reactor is to be used, different zones will generally be used for a continuous process. The pre-reduction step may be carried out at any suitable reaction conditions. However, generally it will be carried out at a lower temperature than that used for the main reaction. The temperature of the pre-reduction step may be from about 150°C to about 250°C and the pressure may be from about 600 to about 1000 psig. The pre-reduction step is found to be particularly useful where the sugar feedstock is an aldose. Whilst not wishing to be bound by any theory it is believed that the terminal aldehyde group of the aldose is reduced and that where the aldose is cyclic, the ring is opened. Some C-C bond cleavage may also occur.

The present invention will now be described with reference to the following examples which are not intended to be limiting on the scope of the invention.

### Examples 1 to 5

These examples demonstrate the effect of varying the reaction temperature in a batch reaction.

0.18g of ruthenium acetylacetonate (from Johnson Matthey), 0.38g of 1,1,1(diphenylphosphino methyl)ethane) (from Aldrich) and tetrahydrofuran (from Aldrich), 20g sorbitol (from Aldrich) and 50g deionised water were weighed into a 300 ml Parr Hastelloy C autoclave which was then sealed. The headspace of the autoclave was purged before being pressurised to approximately 600 psig with hydrogen gas. The stirrer speed was 600 rpm and the reactor heated to the desired temperature. When the temperature was reached, the pressure in the reactor was increased to 1000 psig and the reaction time of 6 hours was considered to have started. The pressure in the autoclave was maintained throughout the reaction by feeding hydrogen gas under regulator control. At the end of the reaction the gas make up was stopped, and the reactor was cooled to room temperature before, the headspace was vented. The liquid products were removed and analysed on a Hewlett Packard HP6890 GC using a J&W 0.32mm,

50m, DB1, with a 1  $\mu$ m phase thickness and using butoxyethanol as an internal standard for quantifying the amounts of propylene glycol, ethylene glycol and glycerol produced.

For the purposes of the results reported below, molar yield is considered to be 100 moles product/moles of feed. Hence if ethylene glycol were the only product a molar yield of 300% could, theoretically be reported for the conversion of sorbitol to products. For polymeric sugars, e.g. starch and sucrose they are considered to have the molecular weight of their monomer units for the molar yield calculation.

The results for various reaction temperatures are set out in Table 1

**Table 1**

Ex No	Temp °C	Ethylene glycol (mol%)	Propylene glycol (mol %)	glycerol (mol %)	Total (Propylene glycol +Ethylene glycol) (mol %)
1	250	48	82	8	130
2	250	50	80	2	130
3	225	51	68	50	119
4	200	57	62	41	119
5	190	42	46	46	88

#### Examples 6 and 7

These examples demonstrate the effect of pressure using a highly volatile solvent.

The method of Examples 1 to 5 was repeated at a temperature of 250°C except that the pressure in the reactor was modulated. The results, which are set out in Table 2, indicate a dramatic loss in selectivity as the pressure is reduced.



**Table 2**

Ex No	Pressure (psig)	Ethylene glycol (mol %)	Propylene glycol (mol %)	Glycerol (mol %)	Total (propylene glycol + ethylene glycol) (mol %)
6	1000	48	82	8	130
7	750	27	27	5	54

**Examples 8 to 13**

This demonstrates that a range of solvents can be employed.

The method of Example 1 was repeated except that the solvent, tetrahydrofuran, was replaced with other solvents in varying amounts.

The results, which are set out in Table 3, illustrate that a range of solvents may be used.

**Table 3**

Ex No	Solvent	Solvent Amount (g)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
8	THF	17.1	48	82	8	130
9	iPA	19.9	34	92	9	126
10	TEGDE	19.0	29	41	<1	70
11	TEGDE	50	56	60	13	116
12	NMP	20.1	7	5	2	12
13	NMP + THF	74.8	104	59	1	163

where THF = tetrahydrofuran, iPA = isopropanol; TEGDE = tetraethyleneglycol dimethylether and NMP = N-methyl pyrrolidone

**Examples 14 to 18**

These examples further demonstrate that a range of solvents may be employed and that their concentration may affect the observed selectivity.

The method of Example 1 was repeated except that the sorbitol was replaced with glucose and the quantity and nature of the solvent and amount of water present were varied.

The results are set out in Table 4.

**Table 4**

Ex No	Solvent	Solvent Amount (g)	Water Amount (g)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
14	THF	20.0	50	30	91	5	121
15	THF	50.0	50	25	55	1	80
16	NMP	20.4	50	20	54	14	74
17	NMP	49.6	50	19	51	1	70
18	NMP	75.0	30	14	34	1	48

**Examples 19 to 24**

These examples demonstrate that the catalyst is suitable for the hydrogenation of a range of sugars as defined in the present invention.

The method of Example 1 was repeated except that the sorbitol was replaced by an alternative substrate.

The results are set out in Table 5. It is postulated that for the given conditions the sorbitol produced a higher yield than the cyclic sugars. Without wishing to be bound by any theory, it is believed that this is due to undesirable reactions occurring while the sugar is in the cyclised state.

**Table 5**

Ex No	Substrate	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
19	Sorbitol	48	82	8	130
24	Starch	31	46	7	77
25	Sucrose	30	67	17	107
26	Glucose	30	91	5	121
27	Xylose	70	43	4	113
28	Arabinose	74	44	5	118

**Examples 25 to 30**

These examples demonstrate the benefits of use of a pre-reduction step.

The method of Example 1 was repeated except that the reaction temperature was initially controlled below the level previously employed for the hydrogenolysis of sugars. The sorbitol was replaced with glucose.

The results are set out in Table 6. It is noted that pre-reduction of the glucose at both 150°C and 200°C improves the selectivity of the reaction such that it is greater than that observed for sorbitol (Example 1). This may be an indication that some hydrogenolysis also takes place at the lower temperature.

**Table 6**

Ex No	Temp1/°C (Time/hrs)	Temp2/°C (Time/hrs)	Temp3/°C (Time/hrs)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
25			250 (6)	30	91	5	121
26		150 (2)	250 (4)	57	90	9	147

27		150 (2)	225 (4)	45	80	22	125
28		200 (2)	250 (2)	58	93	34	151
29		200 (2)	250 (4)	48	94	15	144
30	150 (2)	200 (2)	250 (2)	49	92	19	141

### 5 Examples 31 to 33

These examples further demonstrate the use of a pre-reduction step using N-methyl pyrrolidone as a solvent.

The method of Example 1 was repeated except that the sorbitol was replaced with glucose, the 20g tetrahydrofuran was replaced with 50g N-methyl pyrrolidone and a pre-reduction step was included.

The results are set out in Table 7. Pre-reduction of the glucose at 200°C followed by hydrogenolysis at a higher temperature increases the selectivity towards desirable products. However, increasing the temperature above 260°C appears to have a detrimental effect.

**Table 7**

E.g. No	Temp1/°C (Time/hrs)	Temp2/°C (Time/hrs)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol)
31		250 (6)	19	51	1	70
32	200(2)	260 (4)	63	98	<1	162
33	200(2)	270 (4)	59	50	2	109

### 20 Examples 34 to 38

These examples further illustrate the usefulness of a 'pre-reduction' step in the hydrogenolysis of C<sub>3</sub> alditols.

The method of Example 1 was repeated except that the sorbitol was replaced by xylose or arabinose (C<sub>5</sub> sugars) and a 'pre-reduction' step was employed as outlined below. In Example 38 a mixture of xylose and glucose is used.

The results are set out in Table 8.

**Table 8**

E.g. No	Sugar	Temp1/°C (Time/hrs)	Temp2/°C (Time/hrs)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
34	Xylose		250 (6)	70	43	4	113
35	Arabinose		250 (6)	74	44	5	118
36	Xylose	200 (2)	250 (4)	49	44	1	93
37	Arabinose	200 (2)	250 (4)	79	79	6	158
38	Glucose + Xylose	200 (2)	250 (4)	72	63	10	135

#### Examples 39 to 45

These examples further illustrate the hydrogenolysis of C<sub>5</sub> aldoses using a pre-reduction step and N-methyl pyrrolidone as solvent.

The method of Example 1 was repeated except that the tetrahydrofuran was replaced with 50g of N-methylene pyrrolidone and the sorbitol with xylose.

The results are set out in Table 9. It is noted that in contrast to the results obtained for tetrahydrofuran (Examples 31 to 33), pre-reduction is effective for xylose in N-methyl pyrrolidone. The best results appear to occur with a two hour pre-reduction at 200°C.

Table 9

Ex No	Temp1/°C (Time/hrs)	Temp2/°C (Time/hrs)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (propylene glycol + ethylene glycol) (mol%)
39		260 (6)	50	38	2	88
40		250 (6)	45	47	<1	92
41	200 (2)	260 (4)	79	76	<1	155
42	200 (1)	260 (5)	40	76	<1	116
43	200 (3)	260 (4)	79	39	<1	118
44	200 (2)	260 (2)	77	74	<1	151
45	200 (2)	260 (6)	75	56	1	131

**Examples 48 to 49**

These further demonstrate the suitability of the catalyst for the hydrogenolysis of a range of substrates.

The method of Example 1 was repeated except that the tetrahydrofuran was replaced by 50g of N-methyl pyrrolidone as the solvent, the sorbitol with a range of other substrates and a pre-reduction step was employed. The reaction therefore consisted of 2hrs at 200°C followed by 4hrs at 250°C.

The results are set out in Table 10.

Table 10

Ex No	Substrate	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (propylene glycol + ethylene glycol) (mol%)
46	Glucose	63	98	<1	162

47	Mannose	72	81	8	153
48	Mannitol	77	82	2	159
49	Ribose	80	54	11	134

**Examples 50 to 52**

5 These examples explore the effect of the water concentration.

The method of Examples 39 to 45 was repeated except that glucose was employed as the substrate, and the amounts of water and glucose were modulated as set out in Table 11.

**Table 11**

E.g. No	Water (g)	Glucose (g)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
50	50	20	63	98	<1	162
51	42	28	67	111	1	187
52	20	20	84	70	8	154

**Examples 53 to 55**

15 These examples explore the effect of added base and illustrate that the addition of base does not promote the selectivity of the catalyst as described in other patents. The method of Example 1 was repeated except that an amount of base was added to the reaction. In both cases this caused a small reduction in the amount of desirable products produced. The results are set out in Table 12.

**Table 12**

E.g. No	Additive	Solvent	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
53	None	THF	48	82	8	130

54	NaOH	THF	45	76	2	121
55	NH <sub>4</sub> OH	THF	42	36	1	78

**Examples 56 to 59**

These examples consider the effect of the reaction period and illustrates that the product profile may be varied by varying the reaction period and further illustrates the temperature range over which the catalyst is active.

The method of Example 1 was repeated except that the reaction temperature and reaction period were varied as described in Table 13.

**Table 13**

Ex No	Temp (°C)	Time (hrs)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)	Conversion (wt %)
56	250	6	48	82	8	130	>99
57	250	3	44	76	16	120	>99
58	200	6	46	40	45	86	72
59	150	20	9	9	10	18	>2

**Examples 60 to 63**

These examples demonstrate that with a less volatile solvent the catalyst is relatively insensitive to pressure.

The method of Examples 39 to 45 were repeated except that the reaction pressure was varied. Where sorbitol was employed as a substrate, no 'pre-reduction' step was involved and the total reaction period was 6 hrs. The results are set out in Table 14.



**Table 14**

E.g No	Pressure (psig)	Substrate	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
60	1180	Sorbitol	74	80	3	154
61	1000	Sorbitol	56	67	5	123
62	1213	Glucose	69	81	10	150
63	1000	Glucose	84	70	8	154

**Examples 64 to 71**

These examples illustrate that certain additives can increase the selectivity to the desired product.

The method of Example 1 was repeated, except that an amount of triphenylphosphine was added to the reaction. Where N-methyl pyrrolidone was employed as a solvent, 50g of N-methyl pyrrolidone were used instead of 20g of tetrahydrofuran. The results are set out in Table 15. It can be seen that TPP has a beneficial effect in the presence of certain solvents, notably NMP.

**Table 15**

E.g. No	Additive	Solvent	Pressure (psig)	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
64	None	THF	1000	48	82	8	130
65	TPP	THF	1000	58	72	1	130
66	TPP	THF	1000	51	78	1	129
67	TPP	THF	1000	51	80	2	131
68	TPP	THF	1265	56	67	16	123
69	TPP	NMP	1000	76	76	3	152
70	None	NMP	1000	56	67	<1	123
71	TPP	NMP	1242	68	73	4	141

**Examples 72 to 82**

These examples consider the effect of changing the phosphine and illustrates that tridentate phosphines, in particular facially co-ordinating tripodal phosphines are particularly useful for this reaction. This also provides a comparison with TPP which was employed in the prior art as the ligand of choice.

The method of Example 1 was repeated except that the triphos was replaced by an amount of another ligand as indicated in Table 16.

**Table 16**

Ex No	Ligands (s)	Ligand/Ru ratio	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
72	Triphos	1.2	48	82	8	130
73	Triphos/ TPP	1.2	51	80	2	131
74	Dppe	2.5	10	9	<1	19
75	Dppp	2.6	29	30	1	59
76	TPP	4	4	0.1	2	4
77	None	-	8	2	1	10
78	Dppp	2.6	33	35	9	68
79	Dppp	1.3	8	3	1	11
80	Dppp	4.6	25	27	8	52
81	PCy <sub>3</sub>	7.8	2	0	0	2
82	'Normal' Triphos	1.2	24	35	29	59

"Dppe" is 1,2-bis(diphenylphosphino)ethane, "Dppp" is 1,3-bis(diphenylphosphino)propane, "normal" triphos is 1,1-bis (diphenylphosphinoethyl)phenylphosphine.

**Examples 83 and 84**

A second set of tests were performed using 50g of N-methyl pyrrolidone as a solvent, and at a water loading of 50g. For the Ethphos ligand, the catalyst was pre-formed by heating the ruthenium and phosphine to 200°C for 1hr in the absence of water in N-methyl pyrrolidone.

5 The results are set out in Table 17

**Table 17**

Ex No	Ligands(s)	Ligand/ Ru ratio	Ethylene glycol (mol%)	Propylene glycol (mol%)	Glycerol (mol%)	Total (Propylene glycol + Ethylene glycol) (mol%)
83	Triphos	1.2	48	82	8	130
84	Ethphos	1.0	71	54	19	125

Ethphos is 1, 1, 1-tris (diethylphosphinomethyl) ethane.

**Example 85**

This illustrates that polymeric aldoses such as cellulose will undergo hydrogenolysis in the presence of the catalyst. 11.3g of an NMP solution containing 0.18g of Ru(acac)<sub>3</sub> and 0.38g of triphos (which had been heated to 200°C under nitrogen to coordinate the triphos to the ruthenium), 70g of water and 20g of cellulose (ex Aldrich, 20 micron powder) were loaded into a 300ml hastelloy autoclave. The autoclave was sealed, purged with hydrogen, pressurised to 500psig with Hydrogen and then heated to 200C with stirring. Once 200°C was attained the pressure was increased to 900psig and the reaction started. After 2hrs the reactor was heated to 250°C and the pressure increased to 1000psig. The reaction was left for a further four hours under regulator control. At the end of the reaction 98.3 g of product were recovered containing an orange solution and a solid material (6.1g, unreacted cellulose). The product was analysed by GC using an internal standard. Mol% selectivities EG (52) PG (44). Other products identified in the product mixture by GC-MS include 1-propanol, ethanol, 1-butanol, 1-pentanol, 2-pentanol, 1,2-butanediol and 1,2-pentanediol.